LISTING OF CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in this application.

1-32, (canceled)

33-59. (canceled)

60. (previously presented) An interspinal prosthesis for implantation between a first spinous

process and a second spinal process, the prosthesis comprising:

a first half comprising a coupling portion and a process portion, the coupling portion having a

bore and configured for insertion into the interspinal space between the first spinous process and the

second spinal process, the process portion being sized and configured to be placed on one side of the

first and second spinous processes and being sized and configured to prevent its advancement into the

interspinal space:

a second half comprising a coupling portion and a process portion, the coupling portion

configured to be received within the bore of the coupling portion of the first half, the process portion

being sized and configured to be placed on the other side of the first and second spinous processes and

being sized and configured to prevent advancement into the interspinal space;

a locking mechanism for axially locking the first and second halves together after at least the

coupling portion of the first half has been inserted into the interspinal space;

2

Application No. 10/784,046

Amendment filed December 17, 2007

Response to the Final Office Action dated November 19, 2007

wherein the coupling portion of the first and second halves are sized and configured to be

elastically deformable such that the coupling portion in the area between the first and second spinous

processes has an unstressed diameter and a deformed diameter, said deformed diameter being between

about 10% to about 50% of the unstressed diameter.

61. (previously presented) The interspinal prosthesis of claim 60, the first and second halves

comprising an assembled condition and an unassembled condition, the coupling portions of the first and

second halves insertable into the interspinal space in the unassembled condition, wherein engaging the

coupling portion of the first half with the coupling portion of the second half configures the halves in the

assembled condition.

62. (withdrawn) The interspinal prosthesis of claim 60, wherein the locking mechanism

comprises inner threads on the bore of the first half configured to mate with outer threads on the

coupling portion of the second half such that the first and second halves can be screwed together.

63. (withdrawn) The interspinal prosthesis of claim 62, the locking mechanism further

comprising corresponding engageable ratchet teeth formed on the first and second halves to prevent

unthreading of the first and second halves once the halves have been screwed together.

64. (previously presented) The interspinal prosthesis of claim 60, wherein the coupling portions

of the first and second halves comprise complementary key and keyway surfaces configured to prevent

rotation of the two portions with respect to each other.

3

65. (withdrawn) The interspinal prosthesis of claim 64, wherein the locking mechanism

comprises a nut and bolt combination, the shank of the bolt receivable in complementary bores in the

coupling portions of the first and second halves.

66. (withdrawn) The interspinal prosthesis of claim 60, wherein the locking mechanism

comprises a shoulder in the first half configured to receive a compressible prong on the second half.

67. (withdrawn) The interspinal prosthesis of claim 66, wherein the locking mechanism further

comprises a pin configured to be received within a bore of the compressible prong to render the prong

substantially incompressible.

68. (withdrawn) The interspinal prosthesis of claim 60, wherein the locking member comprises

at least one wire configured to pass through a bore in the recess and projection.

69. (previously presented) The interspinal prosthesis of claim 60, wherein at least a portion of at

least one of the first and second halves is made of an elastomeric material.

70. (previously presented) The interspinal prosthesis of claim 60, wherein at least a portion of at

least one of the first and second halves is made of a metallic material.

71. (previously presented) The interspinal prosthesis of claim 60, wherein at least a portion of at

least one of the first and second halves further comprises a surface for enhancing bone ingrowth.

72. (previously presented) The interspinal prosthesis of claim 71, wherein the surface has a

roughened profile.

4

Application No. 10/784,046

Amendment filed December 17, 2007

Response to the Final Office Action dated November 19, 2007

73. (previously presented) The interspinal prosthesis of claim 71, wherein the surface comprises

a hydroxyapatite coating.

74. (previously presented) The prosthesis of claim 60, the coupling portions configured to

substantially prevent compression of the interspinal space when the coupling portions are inserted in the

interspinal space.

75. (previously presented) The prosthesis of claim 74, the process portions configured to retain

the coupling portions within the interspinal space when the coupling portions are in the locked

configuration.

76. (previously presented) The prosthesis of claim 60, wherein the first half comprises at least

one radially-projecting tab and the second half comprises a groove, at least a portion of the tab

receivable within the groove when the coupling portions are engaged to prevent relative rotational

movement of the first and second halves.

77. (previously presented) The prosthesis of claim 60, wherein the coupling portion of the first

half comprises a stop surface configured to axially engage the second half.

78. (previously presented) The prosthesis of claim 77, wherein stop surface is configured to

separate the process portions of the first and second halves by an amount in the range of from about 2

mm to about 15 mm.

79. (previously presented) The prosthesis of claim 60, wherein the coupling portion of the first

half comprises a cross-sectional dimension of from about 50 mm.sup.2 to about 300 mm.sup.2.

5

Application No. 10/784,046

Amendment filed December 17, 2007

Response to the Final Office Action dated November 19, 2007

80. (previously presented) The prosthesis of claim 79, wherein the process portions of the first

and second halves each have a cross sectional dimension of from about 70 mm.sup.2 to about 500

mm.sup.2.

81.-91 (canceled)